

Solamocta 697 mg/g Powder for Use in Drinking Water for Chickens, Ducks and Turkeys

Authorised

- Amoxicillin trihydrate

Product identification

Medicine name:

Solamocta 697 mg/g Powder for Use in Drinking Water for Chickens, Ducks and Turkeys

Active substance:

Amoxicillin trihydrate

Target species:

Chicken

Turkey

Duck

Route of administration:

In drinking water use

Oral use

Product details

Active substance and strength:

Amoxicillin trihydrate

800.00 milligram(s) / 1.00 gram(s)

Pharmaceutical form:

Powder for use in drinking water

Withdrawal period by route of administration:

In drinking water use:

•

Chicken

- Meat and offal. 1 day

•

Turkey

- Meat and offal. 5 day

•

Duck

- Meat and offal. 9 day

Oral use:

•

Chicken

- Meat and offal. 1 day

•

Turkey

- Meat and offal. 5 day

•

Duck

- Meat and offal. 9 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01CA04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Hungary

Available in:

Hungary

Package description:

1 kg sachet with outside to inside layers of polyethylene terephthalate, aluminum, polyamide, polyethylene (PET/ALU/PA/PE).

100 g sachet with outside to inside layers of polyethylene terephthalate, aluminum, polyamide, polyethylene (PET/ALU/PA/PE).

1 kg sachet with outside to inside layers of polyethylene terephthalate, polyethylene, aluminum, polyethylene (PET/PE/ALU/PE).

500 g sachet with outside to inside layers of polyethylene terephthalate, polyethylene, aluminum, polyethylene (PET/PE/ALU/PE).

250 g sachet with outside to inside layers of polyethylene terephthalate, aluminum, polyamide, polyethylene (PET/ALU/PA/PE).

100 g sachet with outside to inside layers of polyethylene terephthalate, polyethylene, aluminum, polyethylene (PET/PE/ALU/PE).

500 g sachet with outside to inside layers of polyethylene terephthalate, aluminum, polyamide, polyethylene (PET/ALU/PA/PE).

250 g sachet with outside to inside layers of polyethylene terephthalate, polyethylene, aluminum, polyethylene (PET/PE/ALU/PE).

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Eurovet Animal Health B.V.

Marketing authorisation date:

18/12/2015

Manufacturing sites for batch release:

Eurovet Animal Health B.V.

Responsible authority:

Directorate Of Veterinary Medicinal Products

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

18/12/2015

Reference member state:

Netherlands

Procedure number:

NL/V/0288/001

Concerned member states:

Austria Belgium Croatia Czechia Denmark France Germany Greece
Hungary Ireland Italy Latvia Lithuania Luxembourg Poland Portugal
Romania Slovakia Slovenia Spain United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to
www.adrreports.eu/vet